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## INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet 1 of 5

### Complete if Known

Application Number	10/739,208
Filing Date	December 18, 2003
First Named Inventor	KUZMICH, D., et al
Art Unit	1614
Examiner Name	To be Assigned
Attorney Docket Number	9/272

### U. S. PATENT DOCUMENTS

Examiner Initials*	Cite No. <sup>1</sup>	Document Number	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code <sup>2</sup> (if known)			
MS		US- 6,323,199	11/27/2001	Lehmann, M. et al	
MS		US- 5,039,691	08/13/1999	Spagnuolo, C. et al	
MS		US- 6,583,180	06/24/2003	Link, J.T. et al	
MS		US- 6,329,534	12/11/2001	Kym, P.R. et al	
		US- 6,436,986	08/20/2003	Kym, P.R. et al	
		US- 4,880,839	11/14/1999	Tucker, H.	
		US- 2002/0156311	10/24/2002	Link, J.T. et al	
		US- 6,506,766	01/14/2003	Coghlan, M.J. et al	
		US- 6,380,223	04/30/2002	Dow, R.L., et al	
		US- 2003/0232823 A1	12/18/2003	Betageri, R., et al	
		US- 2004/0097574 A1	05/20/2004	Marshall, D.R.	
		US- 2004/0010148 A1	01/15/2004	Kirrane, T.M., Jr. et al	
		US- 2004/0010020 A1	01/15/2004	Kirrane, T.M., Jr., et al	
		US- 2004/0029932 A1	02/12/2004	Bekkali, Y., et al	
MS		US- 2002/0077356 A1	06/20/2002	Jaroch, S., et al	
		US-			
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### FOREIGN PATENT DOCUMENTS

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		Country Code <sup>3</sup> Number <sup>4</sup> Kind Code <sup>5</sup> (if known)				
MS		WO 02/10143	02/07/2002	Schering Aktiengesel.		
		WO 02064550	08/22/2002	Abbott Laboratories		
		WO 99/41256	02/12/1999	Abbott Laboratories		
		WO 02/02565	01/10/2002	Abbott Laboratories		
MS		WO 00/66522	11/09/2000	Pfizer Products, Inc		

Examiner  
Signature

/D Margaret Seaman/

Date  
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Sheet 2 of 5

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MS		EP 0 253 500	02/27/1991	Imperial Chemical Ind		
		EP 0 253 503	12/11/1991	Imperial Chemical Ind		
		GB 2 146 987 A	09/19/1984	Sandoz, Ltd.		
		EP 0 154 528 A2	03/01/1985	Imperial Chemical Ind		
MS		EP 0311447	12/04/1989	Farmos Yhtymä Oy		

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*(Use as many sheets as necessary)*

Sheet	3	of	5
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MS		WO 96/19458	06/27/1996	Ligand Pharmaceut.		
↓		WO 97/27852	08/07/1997	Merck & Company		
		WO 98/54159	12/03/1998	Schering Aktien.		
		WO 00/32584	06/08/2000	Scherin Aktien.		
MS		BE 900594	03/18/1985	Sandoz S.A.		

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		Application Number	10/739,208		
		Filing Date	December 18, 2003		
		First Named Inventor	KUZMICH, D., et al		
		Art Unit	1614		
		Examiner Name	To be Assigned		
Sheet	4	of	5	Attorney Docket Number	9/272

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
MS		Hamann, Lawrence, et al ; Discovery of a potent, Orally active, Nonsteroidal Androgen Receptor Agonist: 4-Ethyl-1,2,3,4-tetrahydro-6-(trifluoromethyl)-8-pyridono[5,6-g]quinoline(LG121071), J. Med Chem, 1999, 42, 210-212	
MS		Pooley, Charlotte, et al; Discovery and Preliminary SAR Studies of a Novel Nonsteroidal Progesterone Receptor Antagonist Pharmacophore, J. Med. Chem 1998, 41, 3461-3466	
MS		Edwards, James, P. et al; 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines as Potent, Orally Active, Nonsteroidal Progesterone Receptor Agonists; The Effect of D-Ring Substituents, J. Med. Chem 1998, 41, 303-310	
MS		Zhi, Lin, et al; 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines: A Novel Class of Nonsteroidal Human Progesterone Receptor Agonists, J. Med. Chem 1998, 41, 291-302	
MS		Zhi, Lin; et al 5-Aryl-1,2,3,4-tetrahydrochromeno[3,4-f]quinolin-3-ones as a Novel Class Of Nonsteroidal Progesterone Receptor Agonists: Effect of A-Ring Modification, J. Med. Chem 1999, 42, 1468-1472	
MS		Tegley, Christopher, et al; 5-Benzylidene 1,2-Dihydrochromeno[3,4-f]quinolines, A Novel Class of Nonsteroidal Human Progesterone Receptor Agonists; J. Med. Chem 1998, 41, 4354-4359	
MS		Edwards, James, P. et al; Preparation, Resolution and Biological Evaluation of 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines as Potent, Orally Active, Nonsteroidal Progesterone Receptor Agonists; J. Med. Chem. 1998, 41, 2779-2785	
MS		Hamann, Lawrence, et al; Synthesis and Biological Activity of a Novel Series of Nonsteroidal, Peripherally Selective androgen Receptor Antagonists Derived from 1,2-Dihydropyridono[5,6-g]quinolines. J. Med. Chem. 1998, 41, 623-639	
		English Translation of WO/02/10143	

Examiner Signature	/D Margaret Seaman/	Date Considered	05/18/2006
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		Filing Date	December 18, 2003
		First Named Inventor	KUZMICH, D., et al
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Sheet	5	of	5
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MS		BEKKALI, Y., et al; Application entitled Glucocorticoid Mimetics, Methods of Making Them, Pharmaceutical Compositions and Uses Thereof, accorded U.S. Serial No. 10/639,131	
MS		CYWIN, C. et al; Application entitled Glucocorticoid Mimetics, Methods of Making Them, Pharmaceutical Compositions and Uses Thereof, accorded U.S. Serial No. 10/785,222	

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		Application Number	To be assigned 10/739208		
		Filing Date	December 18, 2003		
		First Named Inventor	Kuzmich, Daniel et al		
		Art Unit	To be assigned		
		Examiner Name	To be assigned		
Sheet	4	of	4	Attorney Docket Number	9/272

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
MS		Hamann, Lawrence, et al ; Discovery of a potent, Orally active, Nonsteroidal Androgen Receptor Agonist: 4-Ethyl-1,2,3,4-tetrahydro-6-(trifluoromethyl)-8-pyridono[5,6-g]quinoline(LG121071), J. Med Chem, 1999, 42, 210-212	
MS		Pooley, Charlotte, et al; Discovery and Preliminary SAR Studies of a Novel Nonsteroidal Progesterone Receptor Antagonist Pharmacophore, J. Med. Chem 1998, 41, 3461-3466	
MS		Edwards, James, P. et al; 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines as Potent, Orally Active, Nonsteroidal Progesterone Receptor Agonists; The Effect of D-Ring Substituents, J. Med. Chem 1998, 41, 303-310	
MS		Zhi, Lin, et al; 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines: A Novel Class of Nonsteroidal Human Progesterone Receptor Agonists, J. Med. Chem 1998, 41, 291-302	
MS		Zhi, Lin; et al 5-Aryl-1,2,3,4-tetrahydrochromeno[3,4-f]quinolin-3-ones as a Novel Class Of Nonsteroidal Progesterone Receptor Agonists: Effect of A-Ring Modification, J. Med. Chem 1999, 42, 1466-1472	
MS		Tegley, Christopher, et al; 5-Benzylidene 1,2-Dihydrochromeno[3,4-f]quinolines, A Novel Class of Nonsteroidal Human Progesterone Receptor Agonists; J. Med. Chem 1998, 41, 4354-4359	
MS		Edwards, James, P. et al; Preparation, Resolution and Biological Evaluation of 5-Aryl-1,2-dihydro-5H-chromeno[3,4-f]quinolines as Potent, Orally Active, Nonsteroidal Progesterone Receptor Agonists; J. Med. Chem. 1998, 41, 2779-2785	
MS		Hamann, Lawrence, et al; Synthesis and Biological Activity of a Novel Series of Nonsteroidal, Peripherally Selective androgen Receptor Antagonists Derived from 1,2-Dihydropyridono[5,6-g]quinolines. J. Med. Chem. 1998, 41, 623-639	
		<del>English Translation of WO/02/40143</del>	

Examiner Signature	/D Margaret Seaman/	Date Considered	05/18/2006
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